Crystallizing Galactocele – An Unusual Diagnosis on Fine Needle Aspiration Cytology

DHIRAJ B NIKUMBH, SUSHAMA R DESAI, PALLAVI A SHRIGONDEKAR, AJAY BRAHMNALKAR, AVINASH M MANE

A galactocele is a benign cystic lesion that generally occurs during late pregnancy and lactation. Fine Needle Aspiration Cytology (FNAC) reveals a milky fluid, which is diagnostic as well as therapeutic.

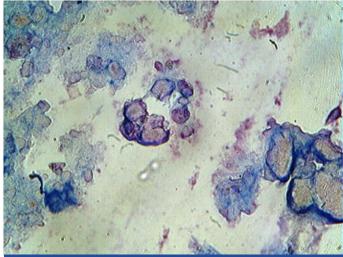
A 27 years old female was referred to our hospital with the chief complaint of a non-tender nodule in her right breast since 2 months. One and a half year back, she had delivered a male baby by a spontaneous, full term, vaginal delivery. She had breast fed her child for one year and thereafter, intermittently to the day of presentation. On physical examination of the breast, a discrete, mobile, nodule measured 1.5 X 1.0 cm was palpable in the upper outer quadrant of the right breast. The clinical impression was of a fibroadenoma. The clinician advised an FNAC of the lesion. We performed 2 needle passes which resulted in the aspiration of a thick, milky material, with a reduction in the size of the lesion. The cytology smears were studied with H and E staining, as well as Leishman's staining. The smears showed numerous, distinct, compact and semitransparent to dark blue/purple crystals on H and E staining [Table/Fig-1] and refractile crystals with defined borders of variable shapes and sizes on Leishman's staining [Table/Fig-2]. The background showed granular, amorphous and proteinaceous

[Table/Fig-1]: Cytological features of crystallizing galactocele as mainly numerous distinct, compact semitransparent to dark blue/purple crystals (H & E,x100).

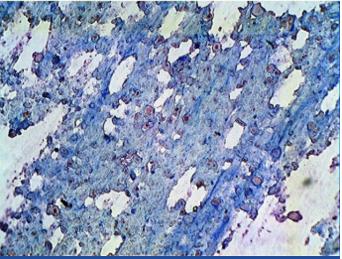
material which was admixed with frothy appearing lipid micelles and variable sized crystals [Table/Fig-3]. The distinct epithelial fragments, bipolar nuclei and fibrous stroma were not seen. In view of the clinical history of the lactation and the cytology, a diagnosis of a crystallizing galactocele was made.

The physiological changes of the breast during pregnancy and lactation make the clinical, radiological and the pathological evaluation of a breast mass particularly challenging [1].

Although most of the palpable breast lesions in the pregnant and lactating patients are benign, 3% of the breast cancers are diagnosed in this population and an evaluation of a breast mass should not be delayed for any reason [2]. FNAC is a well adapted, cost effective technique for the initial evaluation of clinically suspicious breast masses which are found during pregnancy and in the postpartum period [3]. The cytological features of a galactocele which are considered to be specific to pregnancy and lactation, were assessed by only two small series [4] and by a single case report of Raso DS et al., [5].



[Table/Fig-2]: Many refractile blue crystals with defined borders of variable shape and size. (Leishman, x400).



[Table/Fig-3]: Granular, amorphous, proteinaceous material in the background admixed with distinct, variable sized crystals (H&E, x100).

A galactocele is an uncommon benign lesion of the breast and it is defined as an encysted collection of milk products that is lined by a flattened cuboidal epithelium [4, 5]. The major features which are required for the development of a galactocele can be initiated by several features: a secretary breast epithelium, the presence of prolactin stimulation, and some form of ductal obstruction [5]. In addition to lactation, several breast lesions which result in a ductal obstruction or a generalized condition such as a breast surgery, the transplacental passage of prolactin, and oral contraceptives can create the factors for the development of a galactocele [4, 5].

Nevertheless, the diagnosis of a galactocele is often difficult to make on sonography alone, and a pathologic diagnosis is often warranted [4, 5].

To conclude, in a pregnant or a lactating female, a mass which is associated with a milky, viscous aspirate on FNAC, it is of paramount importance, to consider a crystallizing galactocele.

We are reporting this case due to the unusual features of the galactocele, which was diagnosed on FNAC as a crystallizing galactocele. To the best of our knowledge, this is the second case report after Raso DS et al's which was made in 1997.

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AUTHOR(S):

- 1. Dr. Dhiraj B Nikumbh
- 2. Dr. Sushama R Desai
- 3. Dr. Pallavi A Shrigondekar
- 4. Dr. Ajay Brahmnalkar
- 5. Dr. Avinash M Mane

PARTICULARS OF CONTRIBUTORS:

- Department of Pathology, Krishna Institute of Medical Sciences Karad, Maharashtra, India.
- Department of Pathology, Krishna Institute of Medical Sciences Karad, Maharashtra, India.
- 3. Department of Pathology, Krishna Institute of Medical Sciences Karad, Maharashtra, India.
- Department of Medicine, Krishna Institute of Medical Sciences Karad, Maharashtra, India.

5. Department of Pathology, Krishna Institute of Medical Sciences Karad, Maharashtra, India.

NAME, ADDRESS, E-MAIL ID OF THE CORRESPONDING AUTHOR:

Dr. Dhiraj B Nikumbh,

Associate Professor, Department Of Pathology,

JMF's Acpm Medical College, Dhule,

Maharashtra, India.

E-mail: drdhirajnikumbh@rediffmail.com

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